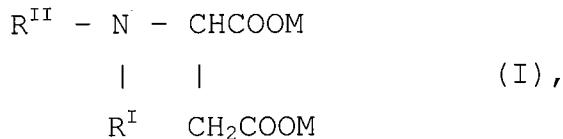


## Amendments to the Claims

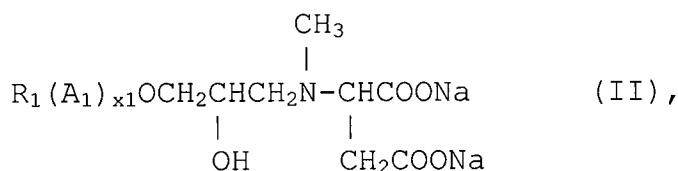
1.(previously presented) A froth flotation process for the enrichment of a calcium phosphate- containing mineral from an ore also containing calcium carbonate, wherein the process is performed in the presence of a collector, wherein said collector is a derivative of aspartic acid of the formula



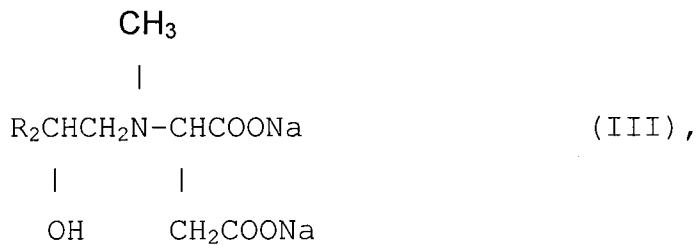
where  $R^I$  is a hydrophobic group containing a hydrocarbon group of 6-24 carbon atoms;  $R^{II}$  is an alkyl group with 1-7 carbon atoms or a group of the formula  $(B)_yH$ , in which B is an alkyleneoxy group with 2-4 carbon atoms and y is a number from 1 to 10; and M is a group selected from the group consisting of a cation or hydrogen.

2.( previously presented) The froth flotation process of claim 1 wherein  $R^I$  is a glycidyl ether group of the formula  $CH_2CH(OH)CH_2O(A_1)_{x1}R_1$ , in which  $R_1$  is a hydrocarbon group with 8-24 carbon atoms,  $A_1$  is an alkyleneoxy group with 2-4 carbon atoms and  $x1$  is a number from 0 to 10; a hydroxyl group of the formula  $CH_2CH(OH)R_2$ , in which  $R_2$  is a hydrocarbon group with 6-22 carbon atoms; a propylene ether group of the formula  $C_3H_6O(A_3)_{x3}R_3$ , in which  $R_3$  is a hydrocarbon group with 8-24 carbon atoms,  $A_3$  is an alkyleneoxy group with 2-4 carbon atoms and  $x3$  is a number from 0-10; or a group of the formula  $R_4$ , where  $R_4$  is a hydrocarbon group containing 8-24 carbon atoms.

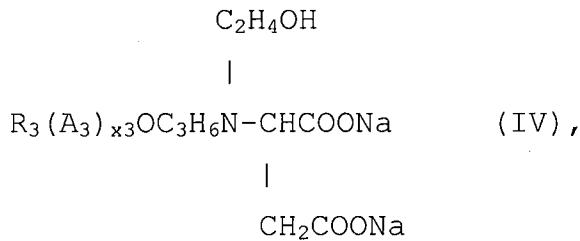
3. (previously presented) The froth flotation process of claim 2, wherein the derivative is selected from the group consisting of



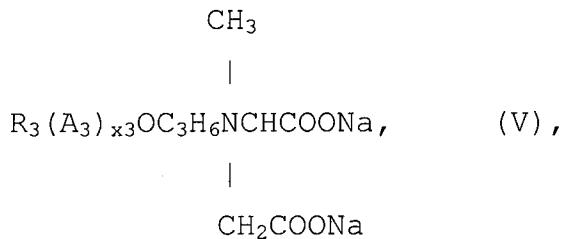
where  $R_1$ ,  $A_1$ ,  $x1$  have the same meanings as in claim 2,



where  $\text{R}_2$  has the same meaning as in claim 2,



where  $\text{R}_3$ ,  $\text{A}_3$  and  $x_3$  have the same meanings as in claim 2, and



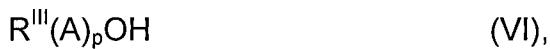
where  $\text{R}_3$ ,  $\text{A}_3$  and  $x_3$  have the same meanings as in claim 2, and mixtures of two or more of the derivatives of formula II, III, IV or V.

4. (previously presented) The froth flotation process of claim 2 wherein  $\text{A}_1$  and  $\text{A}_3$  are both ethyleneoxy and  $x_1$  and  $x_3$  are each independently selected from a number of from 1-4.

5. (previously presented) The froth flotation process of claim 1 wherein  $\text{R}^{\text{II}}$  is methyl, hydroxyethyl or hydroxypropyl.

6. (previously presented) The froth flotation process of claim 1 wherein the derivative is present in an amount of 10-1500 grams per ton of the ore.

7. (previously presented) The froth flotation process of claim 1 wherein the process is performed in the presence of a polar co-collector of the formula

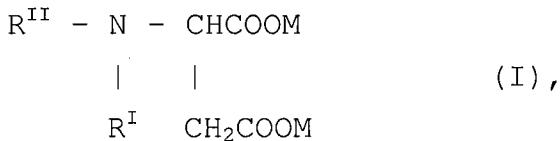


in which  $R^{III}$  is a hydrocarbon group with 8-22 carbon atoms, A is an oxyalkylene group having 2-4 carbon atoms and p is a number from 1-6, or of the formula



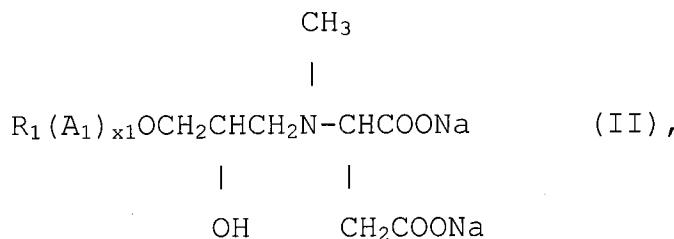
in which  $R^{IV}$  is an aliphatic group having 7-21 carbon atoms, A is an alkyleneoxy group having 2-4 carbon atoms, q is a number from 0-6, and Y is an alkyl group having 1-4 carbon atoms or hydrogen, provided that Y cannot be hydrogen when q is zero.

8. (currently amended) A derivative of aspartic acid of the formula

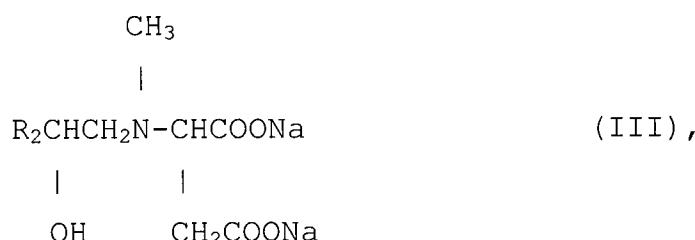


where  $R^I$  is a glycidylether group of the formula  $CH_2CH(OH)CH_2O(A_1)_{x1}R_1$ , in which  $R_1$  is a hydrocarbon group with 8-24 carbon atoms,  $A_1$  is an alkyleneoxy group with 2-4 carbon atoms and  $x1$  is a number from 0 to 10; a hydroxyl group of the formula  $CH_2CH(OH)R_2$ , in which  $R_2$  is a hydrocarbon group with 6-22 carbon atoms; a propylene ether group of the formula  $C_3H_6O(A_3)_{x3}R_3$ , in which  $R_3$  is a hydrocarbon group with 8-24 carbon atoms,  $A_3$  is an alkyleneoxy group with 2-4 carbon atoms and  $x3$  is a number from 0-10; or a group of the formula  $R_4$ , where  $R_4$  is a hydrocarbon group containing 8-24 carbon atoms;  $R^{II}$  is an alkyl group with 1-7 carbons atoms or a group of the formula  $(B)_yH$ , in which B is an alkyleneoxy group with 2-4 carbon atoms and y is a number from 1 to 10; with the proviso that when  $R^{II}$  is an alkyl group with 1-7 carbon atoms then  $R^I$  is not a group  $RCO$ , where R is a C7-C21 alkyl or alkenyl, a group R, where R is a C8-C22 alkyl or alkylene group, or a group  $(CH_2)_3OR$ , where R is a C8-C22 alkyl or alkylene group; wherein when  $R^{II}$  is a group  $-CH_2CH_2OH$  then  $R^I$  is not a group R, where R is a C8-C22 alkyl or alkylene group; and M is a group selected from the group consisting of a cation or hydrogen.

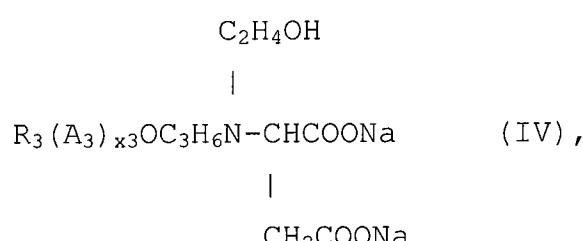
9. (currently amended) The derivative of claim 8, wherein it is selected from the group consisting of



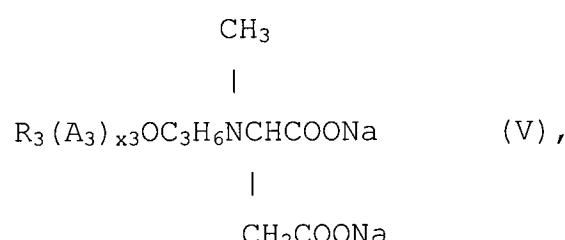
where  $\text{R}_1$ ,  $\text{A}_1$   $x1$  have the same meanings as in claim 82,



where  $\text{R}_2$  has the same meaning as in claim 89,



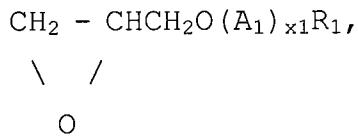
where  $\text{R}_3$ ,  $\text{A}_3$  and  $x3$  have the same meanings as in claim 89, and



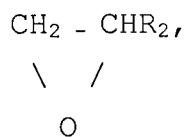
where  $\text{R}_3$ ,  $\text{A}_3$  and  $x3$  have the same meanings as in claim 89, and mixtures of two or more of the derivatives of formula II, III, IV or V.

10. (previously presented) A method of producing a derivative according to claim 8, which comprises reacting maleic acid or a salt thereof under alkaline conditions with

a) a primary amine of the formula  $R''NH_2$ , where  $R''$  has the meaning mentioned above, followed by reacting the intermediate obtained with a glycidylether of the formula



where  $R_1$ ,  $x_1$  and  $\text{A}_1$  have the meanings mentioned above, an epoxide of the formula

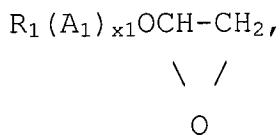


where  $R_2$  has the meaning mentioned above, or a halide compound of the formula

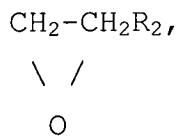
$\text{HalR}_4$ , where  $\text{Hal}$  is a halide and  $R_4$  has the meaning above; or

b) with a primary amine of the formula  $R'NH_2$ , where  $R'$  has the meaning mentioned above, followed by reacting the intermediate obtained with a halide compound of the formula  $\text{HalR}''$ , where  $\text{Hal}$  is a halide and  $R''$  has the meaning mentioned above.

11. (previously presented) The method of claim 10 wherein i) the disodium salt of maleic acid is reacted with N-methylamine and the obtained (N-methyl)aspartate disodium salt is further reacted with a compound of the formula



where  $R_1$ ,  $A_1$  and  $x_1$  have the same meanings as in claim 11 to an aspartate of the formula II, or with a compound of the formula



where  $R_2$  has the same meaning as in claim 2, to obtain an aspartate of the formula III, or

ii) the monosodium salt of maleic acid is reacted with an ether amine of the formula

$R_3(A_3)_{x_3}OC_3H_6NH_2$ ,

where  $R_3$ ,  $A_3$  and  $x_3$  have the meanings mentioned in claim 11 to obtain an intermediate of the formula

$R_3(A_3)_{x_3}OC_3H_6NHCHCOONa$ ,

|

$CH_2COOH$

which intermediate is further reacted with  $Cl(CH_2CH_2O)H$  or  $CH_3Cl$  and with  $NaOH$  to obtain a derivative of formula IV and V, respectively.

12. (previously presented) The froth flotation process of claim 3 wherein  $A_1$  and  $A_3$  is ethyleneoxy and  $x_1$  and  $x_3$  are each independently selected from a number of from 1-4.

13. (previously presented) The froth flotation process of claim 2 wherein  $R^{II}$  is methyl, hydroxyethyl or hydroxypropyl.